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In re Application of:

Belshaw et al.

Serial No: 09/466,568

Filed: December 17, 1999

For: Regulated Gene Therapy Using Specific  
Response Elements

Attorney Docket No. APBI-P16-316

Art Unit: 1636

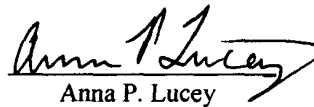
Examiner: Loeb, B. #25

**CERTIFICATE OF MAILING UNDER 37 C.F.R. §1.8(a)**

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**INFORMATION DISCLOSURE STATEMENT IN COMPLIANCE WITH 37 CFR**

**§§ 1.97(b) and 1.98(d)**

Submitted herewith on Form PTO-1449 is a list of publications that applicants and their agents/attorneys have identified during the preparation of this application. In accordance with CFR § 1.98 (d), applicants respectfully submit that *no copy* of any patent, publication, or other information listed on the enclosed Form PTO 1449 is needed because the citations were made in prior application U.S.S.N. 08/605,578, filed February 22, 1996 which is relied upon for an earlier filing date under 35 U.S.C. 120.

Please charge our Deposit Account No. 18-1945 in the amount of \$180.00 covering the fee set forth in 37 CFR 1.17(p).

Applicants respectfully request that the Examiner consider the listed documents and indicate that they were considered by making appropriate notations on the attached Form 1449.

This submission does not represent that a search has been made or that no better art exists. Nor does it constitute an admission that each or all of the listed documents are material or constitute "prior art." If the Examiner applies any of the documents as prior art against any claim in the application and Applicants determine that the cited documents do not constitute "prior art" under United States law, Applicants reserve the right to present to the Office the relevant facts and law regarding the appropriate status of such documents.

Applicants further reserve the right to take appropriate action to establish the patentability of the disclosed invention over the listed documents, should one or more of the documents be applied against the claims of the present application.

Respectfully submitted,  
Ropes & Gray

By: 

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Form PTO-1449		Docket Number (Optional) APBI-P16-316		Application Number 09/466,568	
<b>INFORMATION DISCLOSURE CITATION</b> IN AN APPLICATION (Use several sheets if necessary)		Applicant Belshaw et al.		Group Art Unit 1636	
		Filing Date December 17, 1999			
<b>U.S. PATENT DOCUMENTS</b>					
EXAMINER INITIAL	DOCUMENT NUMBER	DATE	NAME	CLASS	FILING DATE IF APPROPRIATE
AA	5,171,671	12/15/92	Evans et al.		
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<b>FOREIGN PATENT DOCUMENTS</b>					
	DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUBCLASS Translation YES NO
AB	EP 594847	5/4/94	Europe		
AC	WO 93/25533	12/23/93	PCT		
AD	WO 93/23550	11/25/93	PCT		
AE	WO 92/01052	1/23/92	PCT		
<b>OTHER DOCUMENTS</b> <i>(Including Author, Title, Date, Pertinent Pages Etc.)</i>					
AF	Alberg et al. Structure-Based Design of a Cyclophilin-Calcineurin Bridging Ligand. <i>Science</i> 262, 248-250 (1993).				
AG	Albers et al. FKBP, Thought to be Identical to PKCI-2, Does Not Inhibit Protein Kinase C. <i>BioMed. Chem. Lett.</i> 1, 205-210 (1991).				
AH	Albers et al. An FKBP-Rapamycin Sensitive, Cyclin-Dependent Kinase Activity That Correlates with the FKBP Rapamycin-Induced G1 Arrest Point in MG-63 Cells. <i>Annals of N. Y. Acad. Sci.</i> 696, 54-62 (1993).				
AI	Albers et al. The Relationship of FKBP to PKCI-1. <i>Nature</i> 351, 527 (1991).				
AJ	Albers et al. Substrate Specificity for the Human Rotamase FKBP: A View of FK506 and Rapamycin as Leucine (twisted amide)-Proline Mimics. <i>J. Org. Chem.</i> 55, 4984 (1990).				
AK	Andrus et al. Structure-Based Design of an Acyclic Ligand That Bridges FKBP12 and Calcineurin. <i>J. Am. Chem. Soc.</i> 115, 10420-10421 (1993).				
AL	Ben-Levy et al. A oncogenic point mutation confers High Affinity Ligand Binding to the neu Receptor. <i>J. Biol. Chem.</i> 267, 17304-17313 (1992).				
AM	Bergsma et al. The Cyclophilin Multigene Family of Peptidyl-Prolyl Isomerases. <i>J. Biol. Chem.</i> 26, 23204 (1991).				
AN	Bernard et al. High-affinity Interleukin-2 Binding by an Oncogenic Hybrid Interleukin-2 Epidermal Growth Factor Receptor Molecule. <i>PNAS</i> 84, 2125-2129 (1987).				
AO	Bierer et al. The Effect of the Immunosuppressant FK506 on Alternate Pathways of T Cell Activation. <i>Eur. J. Immunol.</i> 21, 439-445 (1991).				

Part of Paper No. 25

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		Filing Date December 17, 1999			
	AP	Bierer et al. Mechanisms of Immunosuppression by FK506: Preservation of T Cell Transmembrane Signal Transduction. <i>Transplantation</i> 49, 1168 (1990).			
	AQ	Bierer et al. Probing Immunosuppressant Action with a Nonnatural Immunosuppressive Ligand. <i>Science</i> 250, 556 (1990).			
	AR	Bierer et al. Two Distinct Signal Transmission Pathways in T Lymphocytes are Inhibited by Complexes Formed Between an Immunophilin and Either FK506 or Rapamycin. <i>PNAS</i> 87, 9231 (1990).			
COPY OF PAPERS ORIGINALLY FILED	AS	Bonnerot et al. Role of associated $\gamma$ -chain in Tyrosine Kinase Activation via Murine FcRIII. <i>EMBO J.</i> 11, 2747-2757 (1992).			
	AT	Bram et al. Identification of the Immunophilins Capable of Mediating Inhibition of Signal Transduction by Cyclosporin A and FK506: Roles of Calcineurin Binding and Cellular Location. <i>Mol. Cell. Biol.</i> 13, 4760-4769 (1993).			
	AU	Byrn et al. Biological Properties of a CD4 Immunoaderhesin. <i>Nature</i> 344, 667-670 (1990).			
	AV	Cantley et al. Oncogenes and signal transduction. <i>Cell</i> 64, 281-302 (1991).			
	AW	Chan et al. The $\zeta$ Chain is associated with a Tyrosine Kinase and upon T-Cell Antigen Receptor Stimulation Associates with ZAP-70, a 70-kDa Tyrosine Phosphoprotein. <i>PNAS</i> 88, 9166-9170 (1991).			
	AX	Chung et al. Rapamycin-FKBP specifically blocks growth-dependent activation of and signaling by the 70 kd S6 protein kinases. <i>Cell</i> 69, 1227 (1992).			
	AY	Clark et al. The B Cell Antigen Receptor Complex: Association of Ig- $\alpha$ and Ig- $\beta$ with Distinct Cytoplasmic Effectors. <i>Science</i> 258, 123-126 (1992).			
	AZ	Clipstone et al. Calcineurin: Molecular analysis of its interaction with drug-immunophilin complexes and its role in the regulation of NF-AT. <i>J. Cell. Biochem. Suppl.</i> 0 (18B) 274, Abstract #1410 (1994).			
	BA	Crabtree, G. R. IL-2 receptor in the pathogenesis of human lymphoma. Abstract of NIH Grant R01CA39612 (1987).			
	BB	Crabtree, G. R. Pathways of T lymphocyte activation. Abstract of NIH Grant R01CA39612 (1991).			
	BC	Crabtree, G. R. Pathways of T lymphocyte activation. Abstract of NIH Grant R01CA39612 (1988).			
	BD	DiLella et al. Chromosomal Band Assignments of the Genes Encoding Human FKBP12 and FKBP13. <i>Biochem. Biophys. Res. Commun.</i> 189, 819-823 (1992).			
	BE	Donald et al. C10 N-Acyl Modified FK-506: A Possible Hybrid Analogue of the Transition State of Petidyl-Prolyl Cis-Trans Isomerization. <i>Tetrahedron Letters</i> 31, 1375-1378 (1991).			
	BF	Durand. Characterization of Antigen Receptor Response Elements within the Interleukin-2 Enhancer. <i>Mol. Cell. Biol.</i> 8, 1715 (1988).			

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		Filing Date December 17, 1999	Group Art Unit 1636
	BG	Eberle & Juninger. Synthesis of the Main Metabolite (OL-17) of Cyclosporin A. <i>J. Org. Chem.</i> 57, 2689-2691 (1992).	
	BH	Edalji et al. High-Level Expression of Recombinant Human FK-Binding Protein from a Fusion Precursor. <i>J. Prot. Chem.</i> 11, 213 (1992).	
	BI	Eiseman et al. Signal Transduction by the Cytoplasmic Domains of FcεRI-γ and TCR0J-γ in Rat Basophilic Leukemia Cells. <i>J. Biol. Chem.</i> 267, 21027-21032 (1992).	
	BJ	Emmel et al. Cyclosporin A Specifically Inhibits Function of Nuclear Proteins Involved in T-Cell Activation. <i>Science</i> (1989).	
	BK	Engel et al. High-Efficiency Expression and Solubilization of Functional T-Cell antigen Receptor Heterodimers. <i>Science</i> 256, 1318-1321 (1992).	
	BL	Evans et al. Mechanistic Study of the Rhodium(I)- and Iridium(I)- Catalyzed Hydroboration Reactions: Scope and Synthetic Applications. <i>J. Am. Chem. Soc.</i> 114, 6671-6679 (1992).	
	BM	Fields, S. & Song, O. A Novel Genetic System to Detect Protein-Protein Interactions. <i>Nature</i> 340, 245-246 (1989).	
	BN	Fischer et al. Mip protein of <i>Legionella pneumophila</i> exhibits peptidyl-prolyl-cis/trans isomerase (Pplase) activity. <i>Mol. Microbiol.</i> 6, 1375 (1992).	
	BO	Fisher et al. On the Remarkable Propensity for Carbon-Carbon Bond Cleavage Reactions in the C8-C10 Region of FK-506. <i>J. Org. Chem.</i> 56, 2900-2907 (1991).	
	BP	Flanagan et al. Intracellular signal transmission: a novel role for the prolyl isomerases. <i>J. Cell. Biochem. Suppl.</i> 0 (16 Part A) 61, Abstract #B005 (1992).	
	BQ	Flanagan et al. Nuclear Association of a T-Cell Transcription Factor Blocked by a Tyrosine Factor Blocked by FK-506 and Cyclosporin A. <i>Nature</i> 352, 803-807 (1991).	
	BR	Flanagan et al. Nuclear association of a transcription factor essential for T cell activation by cyclosporin A and FK506. <i>J. Cell. Biochem. Suppl.</i> 0 (16 Part B), 237, Abstract #H514 (1992).	
	BS	Francavilla et al. Inhibition of Liver, Kidney, and Intestine Regeneration by Rapamycin. <i>Transplantation</i> 53, 496-498 (1992).	
	BT	Fretz et al. Rapamycin and FK506 Binding Proteins (Immunophilins). <i>J. Am. Chem. Soc.</i> 113, 1409 (1991).	
	BU	Friedman & Weissman. Two Cytoplasmic Candidates for Immunophilin Action are Revealed by Affinity for a New Cyclophilin: One in the Presence and One in the Presence and One in the Absence of CsA. <i>Cell</i> 66, 799 (1991).	
	BV	Fuh et al. Rational design of potent antagonists to the human growth hormone receptor. <i>Science</i> 256, 1677-1680 (1992).	
	BW	Galat et al. A Rapamycin-Selective 25 kDa Immunophilin. <i>Biochemistry</i> 31, 2427-2434 (1992).	

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	BX	Ghosh et al. N,N'-Dimethylimidyl Carbonate: A Useful Reagent for Alkoxyacylation of Amines. <i>Tetrahedron Letters</i> 33,2781-2784 (1992).	
	BY	Gottschalk et al. The Carboxy Terminal 100 Amino Acid Portion of the Insulin Receptor is Important for Insulin Signaling to Pyruvate Dehydrogenase. <i>Biochem. Biophys. Res. Comm.</i> 189, 906-911 (1992).	
	BZ	Haendler et al. Yeast cyclophilin: isolation and characterization of the protein, cDNA and gene. <i>EMBO J.</i> 6, 947 (1987).	
	CA	Haendler et al. Yeast cyclophilin: isolation and characterization of the protein, cDNA and gene. <i>Gene</i> 83, 39 (1989).	
	CB	Harding et al. A Receptor for the Immunosuppressant FK506 is a cis-trans Peptidyl-Prolyl Isomerase. <i>Nature</i> 341, 758 (1989).	
	CC	Herbst et al. Substrate Phosphorylation Specificity of the Human c-kit Receptor Tyrosine Kinase. <i>J. Biol. Chem.</i> 266, 19908-19916 (1991).	
	CD	Howard et al. The CD3 $\zeta$ Cytoplasmic Domain Mediates CD2-Induced T Cell Activation. <i>J. Exp. Med.</i> 176, 139-145 (1992).	
	CE	Hultsch et al. Immunophilin Ligands Demonstrate Common Features of Signal Transduction Leading to Exocytosis or Transcription. <i>PNAS</i> 88, 6229-6233 (1991).	
	CF	Hultsch et al. Inhibition of IgE Receptor-Mediated Exocytosis from Rat Basophilic Leukemia Cells by FK506 is Reversed by Rapamycin: Evidence for Common Signaling Pathways in Mast Cells and T Lymphocytes. <i>FASEB J.</i> 5, A1008 [3705] (1991).	
	CG	Hung & Schreiber. CDNA Cloning of a Human 25 kDa FK506 and Rapamycin Binding Protein. <i>Biochem. Biophys. Res. Comm.</i> 184, 733 (1992).	
	CH	Ikeda et al. Structural Basis for Peptidomimicry by a Natural Product. <i>J. Am. Chem. Soc.</i> 116, 4143-4144 (1994).	
	CI	Irving & Weiss. The Cytoplasmic Domain of the T Cell Receptor $\zeta$ Chain is Sufficient to Couple to Receptor-Associated Signal Transduction Pathways. <i>Cell</i> 64, 891-901 (1991).	
	CJ	Itoh & Nagata. A Novel Protein Domain Required for Apoptosis. <i>J. B. C.</i> 268, 10932 (1993).	
	CK	Itoh et al. Effect of bcl-2 on Fas Antigen Mediated Cell Death. <i>J. Immunol.</i> 151, 621-627 (1993).	
	CL	Jin et al. Molecular cloning of a membrane-associated human FK506- and rapamycin-binding protein, FKBP-13. <i>PNAS</i> 88, 6677 (1991).	
	CM	Kao et al. Nuclear target of cyclosporin A and FK506 action is specifically bound by a heterodimeric protein comprising molecular weights 90K and 45K. <i>J. Cell. Biochem. Suppl.</i> 0 (16 Part B), 239, Abstract #H523 (1992).	
	CN	Kaye et al. Effects of Cyclosporin A and FK506 on Fce Receptor type I-Initiated Increases in Cytokine mRNA in Mouse Bone Marrow-Derived Progenitor Mast Cells: Resistance to FK506 is Associated with a Deficiency in FKBP12. <i>PNAS</i> 89, 8542-4546 (1992).	

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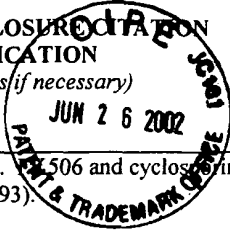
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	CO	Ke et al. Crystal Structures of Cyclophilin A Complexed with Cyclosporin A and N-methyl-4-[(E)-2-Butenyl]-4,4-Dimethyltheanine Cyclosporin A. <i>Structure</i> 2, 33-44 (1994).	
	CP	Kinet. Antibody-Cell Interactions: Fc Receptors. <i>Cell</i> 57, 351-354 (1989).	
	CQ	Krishnamurthy. Lithium Tris[(3-ethyl-3-pentyl)oxy]aluminum Hydride. A New Remarkably Chemoselective Reagent for the Reduction of Aldehydes in the Presence of Ketones. <i>J. Org. Chem.</i> 46, 4628-4629 (1981).	
	CR	Kruskal et al. Phagocytic Chimeric Receptors Require Both Transmembrane and Cytoplasmic Domains from the Mannos Receptor. <i>J. Exp. Med.</i> 176, 1673-1680 (1992).	
	CS	Lammers et al. Differential Signaling Potential in Insulin- and IGF-1-receptor Cytoplasmic Domains. <i>EMBO J.</i> 8, 1369-1375 (1989).	
	CT	Lane et al. Complete Amino Acid Sequence of the FK506 and Rapamycin Binding Protein, FKBP, Isolated from Calf Thymus. <i>J. Prot. Chem.</i> 10, 151-160 (1991).	
	CU	Lanier et al. Co-association of CD3 $\zeta$ with a Receptor (CD16) for IgG Fc on Human Natural Killer Cells. <i>Nature</i> 342, 803-805 (1989).	
	CV	Larson & Nuss. Cyclophilin-dependent stimulation of transcription by cyclosporin A. <i>PNAS</i> 90, 148 (1993).	
	CW	Lee et al. Functional Dissection of Structural Domains in the Receptor for Colony Stimulating Factor-1. <i>J. Biol. Chem.</i> 267, 16472-16483 (1992).	
	CX	Lee et al. HER2 Cytoplasmic Domain Generates Normal Mitogenic and Transforming Signals in a Chimeric Receptor. <i>EMBO J.</i> 8, 167-173 (1989).	
	CY	Lehtola et al. A chimeric EGFR/neu receptor in functional analysis of the neu oncoprotein. <i>Acta Oncologia</i> 31, 147-150 (1992).	
	CZ	Lehtola et al. Receptor Downregulation and DNA Synthesis are Modulated by EGF and TPA in Cells Expressing an EGF/neu Chimera. <i>Growth Factors</i> 1, 323-334 (1989). - ABSTRACT ONLY	
	DA	Lehvaslaiho et al. A Chimeric EGF-R-neu Proto-Oncogene Allows EGF to Regulate neu Tyrosine Kinase and Cell Transformation. <i>EMBO J.</i> 8, 159-166 (1989).	
	DB	Letourner & Klausner. Activation of T Cells by a Tyrosine Kinase Activation Domain in the Cytoplasmic Tail of CD3 $\epsilon$ . <i>Science</i> 258, 123-126 (1992).	
	DC	Lev et al. Receptor functions and ligand-dependent transforming potential of a chimeric kit proto-oncogene. <i>Mol. Cell. Biol.</i> 10, 6064-6068 (1990).	
	DD	Lev et al. A Specific Combination of Substrates is Involved in Signal Transduction by the kit-Encoded Receptor. <i>EMBO J.</i> 10, 647-654 (1991).	
	DE	Levaslaiho et al. Regulation by EGF is maintained in an overexpressed chimeric EDG/neu receptor tyrosine kinase. <i>Biochem.</i> 42, 123-133 (1990).	

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	DF	Liu. FK506 and cyclosporin, molecular probes for studying intracellular signal transduction. <i>Immunology Today</i> 14, 290 (1993).	
	DG	Liu et al. Calcineurin is a Common Target of Cyclophilin-Cyclosporin A and FKBP-FK506 Complexes. <i>Cell</i> 66, 807 (1991).	
	DH	Liu et al. Cloning, expression, and purification of human cyclophilin in Escherichia coli and assessment of the catalytic role of cysteines by site-directed mutagenesis. <i>PNAS</i> 87, 2304 (1990).	
	DI	Liu et al. Inhibition of T Cell Signaling by Immunophilin-Ligand Complexes Correlates With Loss of Calcineurin Phosphatase Activity. <i>Biochemistry</i> 31, 3896-3901 (1992).	
	DJ	Maki et al. Complementary DNA encoding the human T-cell FK506-binding protein, a peptidylprolyl cis-trans isomerase distinct from cyclophilin. <i>PNAS</i> 87, 5440 (1990).	
	DK	Mares et al. A Chimera between Platelet-Derived Growth Factor B-receptor and Fibroblast Growth Factor Receptor-1 Stimulates Pancreatic $\beta$ -DNA Synthesis in the Presence of PDGF-BB. <i>Growth Factors</i> 6, 93-101 (1992).	
	DL	Margolis et al. All Autophosphorylation Sites of Epidermal Growth Factor (EGF) Receptor and HER2/neu are Located in their Carboxyl-Terminals Tails. <i>J. Biol. Chem.</i> 264, 10667-10671 (1989).	
	DM	Mattila et al. The Actions of Cyclosporin A and FK506 Suggest A Novel Step in the Activation of T Lymphocytes. <i>EMBO J.</i> 9, 4425-4433 (1990).	
	DN	Meyer et al. Synthetic Investigations of Rapamycin. 1. Synthesis of a C10-C21 Fragment. <i>J. Org. Chem.</i> 57, 5058-5060 (1992).	
	DO	Michnick et al. Solution Structure of FKBP, a Rotamase Enzyme and Receptor for FK506 and Rapamycin. <i>Science</i> 252, 836-839 (1991).	
	DP	Moe et al. Transmembrane Signaling by a Chimera of the Escherichia coli Aspartate Receptor and the Human Insulin Receptor. <i>PNAS</i> 86, 5683-5687 (1989).	
	DQ	Nakatsuka et al. Total Syntheses of FK506 and an FKBP Probe Reagent, (C8, C9-13C2)-FK506. <i>J. Am. Chem. Soc.</i> 112, 5583 (1990).	
	DR	Nussbaumer et al. C9-Imino and C10-Amino Derivatives of Ascomycin (21-Ethyl-FK506). <i>Tetrahedron Letters</i> 33, 3845-3846 (1992).	
	DS	Orloff et al. Family of Disulphide-Linked Dimers Containing the $\zeta$ and $\eta$ Chains of the T-Cell Receptor and the $\gamma$ Chain of the Fc Receptors. <i>Nature</i> 347, 189-191 (1990).	
	DT	Palmiter et al. Transgenic Mice. <i>Cell</i> 41, 343-345 (1985).	
	DU	Patchett et al. Analogs of Cyclosporin A Modified at the D-ALA <sup>8</sup> Position. <i>J. Antibiotics</i> 45, 94-102 (1992).	
	DV	Peles et al. Regulated Coupling of the Neu Receptor to Phosphatidylinositol. <i>J. Biol. Chem.</i> 267, 12266-12274 (1992).	

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	DW	Price et al. Immunophilin B: A second cyclophilin gene encodes a peptidyl-prolyl isomerase with a signal sequence. <i>PNAS</i> 88, 1903 (1991).
	DX	Ptashne et al. Activators and Targets. <i>Nature</i> 346, 329-331 (1990).
	DY	Ragan et al. Studies of the Immunosuppressive Agent FK506: Synthesis of an Advanced Intermediate. <i>J. Org. Chem.</i> 54, 4267 (1989).
	DZ	Reins et al. Anti-epidermal growth factor receptor monoclonal antibodies affecting signal transduction. <i>J. Cell. Biol.</i> 51, 236-248 (1993).
	EA	Riedel et al. Cytoplasmic Domains Determine Signal Specificity, Cellular Routing Characteristics and Influence Ligand Binding of Epidermal Growth Factor and Insulin Receptors. <i>EMBO J.</i> 8, 2943-2954 (1989).
	EB	Romeo et al. Cellular immunity to HIV activated by CD4 fused to T cell or Fc receptor polypeptides. <i>Cell</i> 64, 1037-1046 (1991).
	EC	Romo et al. Synthetic Investigations of Rapamycin. 2. Synthesis of a C22-C42 Fragment. <i>J. Org. Chem.</i> 57, 5060-5063 (1992).
	ED	Romo et al. Total Synthesis of Rapamycin Using an Evans-Tischenko Fragment Coupling. <i>J. Am. Chem. Soc.</i> 115, 7906-7907 (1993).
	EE	Rosen, M. K. The molecular basis of receptor-ligand-receptor interactions: Studies of the immunophilin FKBP12, Abstract of Doctoral Thesis (1993).
	EF	Rosen et al. Activation of an Inactive Immunophilin by Mutagenesis. <i>J. Am. Chem. Soc.</i> 115, 821-822 (1993).
	EG	Rosen et al. Inhibition of FKBP Rotamase Activity by Immunosuppressant FK506: A Twisted Amide Surrogate. <i>Science</i> 248, 863 (1990).
	EH	Rosen et al. Natural Products as Probes of Cellular Function: Studies of Immunophilins. <i>Angew. Chemie. Int. Ed. Eng.</i> 31, 384-400 (1992).
	EI	Rosen et al. Proton and Nitrogen Sequential Assignments and Secondary Structure Determination of the Human FK506 and Rapamycin Binding Protein. <i>Biochemistry</i> 30, 4774-4789 (1991).
	EJ	Rosen et al. Study of Receptor-Ligand Interactions Through Receptor Labeling and Isotope-Edited NMR. <i>J. Org. Chem.</i> 56, 6262 (1991).
	EK	Roussel et al. Antibody-Induced Mitogenicity Mediated by a Chimeric CD2-c-fms Receptor. <i>Mol. Cell. Biol.</i> 10, 2407-2412 (1990).
	EL	Rudert et al. Apoptosis in L929 cells expressing a CD40/Fas chimeric receptor: Dissociation of stimulatory from inhibitory death signaling functions. <i>Biochem. Biophys. Res. Comm.</i> 204, 1102 (1994).
	EM	Sampson & Gotschlich. Neisseria meningitidis encodes an FK506-inhibitable rotamase. <i>PNAS</i> 89, 1164 (1992).

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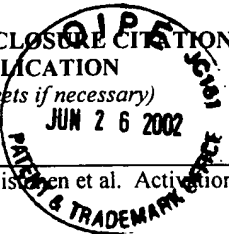
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EY	Schultz et al. Atomic Structure of the Immunophilin FKBP13-FK506 Complex: Insights Into the Composite Binding Surface for Calcineurin. <i>J. Am. Chem. Soc.</i> 116, 3129-3130 (1994).		
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		Applicant Belshaw et al.	
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	FE	Sistonen et al. Activation of the neu Tyrosine Kinase. <i>J. Cell. Biol.</i> 109, 1911-1919 (1989).	
	FF	Smith et al. FKBP54, a Novel FK506 Binding Protein in Avian Progesterone Receptor Complexes and HeLa Extracts. <i>J. Biol. Chem.</i> 268, 24270-24273 (1993).	
	FG	Somers et al. Synthesis and Analysis of 506BD, a High Affinity Ligand to the Immunophilin, FKBP. <i>J. Am. Chem. Soc.</i> 113, 8045-8056 (1991).	
	FH	Standaert, R. F. Biochemical and structural studies of the FK506- and rapamycin-binding proteins (FKBPs). <i>Abstract of Doctoral Thesis</i> (1992).	
	FI	Standaert et al. Molecular cloning and overexpression of the human FK506-binding protein FKBP. <i>Nature</i> 346, 671 (1990).	
	FJ	Tai et al. Association of a 59-Kilodalton Immunophilin with the Glucocorticoid Receptor Complex. <i>Science</i> 256, 1315-1318 (1992).	
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	FL	Tanida et al. Yeast Cyclophilin-related gene encodes a nonessential second peptidyl-prolyl cis-trans isomerase with the secretory pathway. <i>Transplantation Proceedings</i> 23, 2856 (1991).	
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